WHAT IS CLAIMED IS:

- A nitrosated and/or nitrosylated phosphodiesterase inhibitor having the formula NO₀-PDE wherein is 1 or 2.
- The nitrosated and/or nitrosylated phosphodiesterase inhibitor of claim 1 which
 is nitrosylated or nitrosated through an oxygen, sulfur, carbon or nitrogen site on the
 phosphodiesterase inhibitor.
- The nitrosated and/or nitrosylated phosphodiesterase inhibitor of claim 1 which is selected from the group consisting of:
 - (I) compounds having the structure:

wherein,

R1 is alkoxy, cycloalkoxy, halogen, or

R₂ is hydrogen, alkoxy, or haloalkoxy; and

R₃ is selected from:

(i)

R₄—N

(ii) R₄—N

(iv)

-68-

(iii) S N N N

D-0-{N-N

R₅ N S R₄

wherein

D is selected from (i) -NO; (ii) -NO₂; (iii) -C(R_d)-O-C(O)-Y-Z-[C(R_e)(R_f)]_p-T-Q in which R_d is hydrogen, lower alkyl, cycloalkyl, aryl, alkylaryl, aryl or heteroaryl, Y is oxygen, sulfur, or NR_i in which R_i is hydrogen, lower alkyl, R_e and R_f at each occurrence are independently selected from hydrogen, lower alkyl, cycloalkyl, aryl, heteroaryl, arylalkyl, amino, alkylamino, amido, alkylamido, dialkylamino, carboxy, or taken together are carbonyl, cycloalkyl or bridged cycloalkyl, p is an integer from 1 to 6, T is a covalent bond, oxygen, sulfur or nitrogen, Z is selected from a covalent bond, alkyl, cycloalkyl, aryl, heteroaryl, arylalkyl or arylheterocyclic ring, and Q is selected from -NO or -NO₂; (iv) -C(O)-T¹-Z-[C(R_e)(R_f)]_p-T²-Q wherein T¹ and T² are independently selected from T and R_e, R₆ p, Q, Z, and T are as defined in this specification; (v) -C(O)-T₁-C(C_R-((R_f))_p-T²-Q)_p wherein G is (i) a covalent bond; (ii) -T-C(O)-; (iii) -C(O)-T₁ or (iv) Y, and wherein R_e, R₆ p, Q, T, Y, and Z are as defined in this specification; (v) -C(O)-T[C(R_e)(R_f)]_p, T²-Q wherein R_y and R_z are independently selected from -T¹-[C(R_e)(R_f)]_p-T²-Q wherein G, R_e, R₆ p, Q, T, T¹, and T² are as defined in this specification;

 R_4 is selected from (i) hydrogen, (ii) -C(R_d)-O-C(O)-Y-Z-[C(R_e)(R_i)] $_p$ -T-Q, (iii) -C(O)-T¹-[C(R_e)(R_i)] $_p$ -T²-Q, (iv) -C(O)-Z-[G-[C(R_e)(R_i)] $_p$ -T-Q] $_p$; and wherein R_d , R_e , R_f , p, G, T, T^1 , T^2 , Q, Y, and Z are defined as in this specification;

 R_5 is selected from a lone pair of electrons or ${}^{-}$ C(R_d)-O-C(O)-Y-Z-[C(R_c)(R_d)] $_p$ -T-Q wherein R_d , R_e , R_f , p, T, T^1 , T^2 , Q, Y, and Z are defined as in this specification;

 R_{11} and R_{12} are independently selected from hydrogen or R_4 wherein R_4 is as defined in this specification with the provision that R_{11} and R_{12} are not both hydrogen;

X is a halogen and;

D₁ is selected from D or hydrogen and wherein D is as defined in this specification.

(II) compounds having the structure:

II

wherein,

R4 is as defined in this specification;

R₈ is selected from hydrogen or lower alkyl;

Rq is selected from hydrogen or halogen; and

R₁₀ is selected from:

(i) hydrogen

wherein R₈ is as defined in this specification.

(III) compounds having the structure:

Ш

wherein,

E is selected from nitrogen or -CH-;

G is selected from nitrogen or -C(R₈)-;

R₂₁ is selected from:

(ii)

R₂₂ is selected from R₁₂ or lower alkyl; and

 R_8 , R_{11} , and R_{12} are as defined in this specification.

(IV) compounds having the structure:

IV

wherein,

F is selected from -CH₂- or sulfur;

 R_{4} and R_{8} are as defined in this specification; and

R₁₃ is selected from:

(ii)
$$H_3CO$$
 H_3CO
 H_3CO

(vii)

R₆

N

CH₃

wherein,

 R_6 and R_7 are independently selected from hydrogen or R_4 wherein R_4 is as defined in this specification.

wherein,

The first first time and the first of the fi

 R_4 is as defined in this specification; and R_{14} is selected from:

(i) (ii)

wherein R₆ is as defined in this specification.

(VI) compounds having the structure:

VI

wherein.

R₁₅ is hydrogen, lower alkyl, R₄, or -(CH₂)₄-C(CH₃)₂-O-D₁;

R₁₆ is lower alkyl; and

 R_{17} is hydrogen, lower alkyl, CH₃-C(O)-CH₂-, CH₃-O-CH₂-, or D with the provision that either R_{15} or R_{17} must be selected to contain D and wherein D and D₁ are as defined in this specification.

(VII) compounds having the structure:

VII

wherein,

 $R_{\mbox{\tiny 4}}$ and $R_{\mbox{\tiny 8}}$ are as defined in this specification and

R₁₈ is selected from:

and wherein R₈ is as defined in this specification.

(VIII) compounds having the structure:

VIII

wherein,

R₁₉ is selected from:

(i)

and wherein R_4 , R_{11} , and R_{12} are defined as in this specification.

(IX) compounds having the structure:

(ii)

(iv)

wherein,

R₂₀ is selected from:

(ii)

and wherein R4 is defined as in this specification.

(X) compounds having the structure:

wherein,

a is an integer from 2 to 3 and D and D₁ are defined as in this specification.

(XI) compounds having the structure:

wherein D and D1 are defined as in this specification.

(XII) compounds having the structure:

ХII

wherein,

J is selected from:

K is selected from:

(ii)

wherein V is carbon or nitrogen;

 R_{23} , R_{24} , R_{25} , R_{26} , R_{27} , R_{28} , R_{29} , and R_{30} are independently selected from hydrogen, halogen, alkoxy, nitrile, carboxamido, or carboxyl; and wherein p, R_{5} , R_{5} , T, T^{1} , T^{2} , Y and D are defined as in this specification.

(XIII) compounds having the structure:

wherein.

 R_{31} is alkyl, halogen, haloalkyl, or haloalkoxy; R_{32} is selected from D_1 or -C(O)- R_8 ; and

wherein D1 and R8 are defined as in this specification.

- 4. A composition comprising a therapeutically effective amount of the phosphodiesterase inhibitor of claim 1 and a one to ten fold molar excess of a compound that donates, transfers or releases nitrogen monoxide as a charged species, i.e., nitrosonium (NO*), or nitroxyl (NO*), or as the neutral species, nitric oxide (NO*)or induces the production of endogenous EDRF and a pharmaceutically acceptable carrier.
- A method for treating male impotence in humans which comprises administering to an individual in need thereof a therapeutically effective amount of a nitrosated or nitrosylated PDE inhibitor of claim 1.
- 6. A method for treating female sexual dysfunction in humans which comprises administering to an individual in need thereof a therapeutically effective amount of a nitrosated or nitrosylated PDE inhibitor of claim 1.
- A method for treating anal disease in humans which comprises administering to an individual in need thereof a therapeutically effective amount of a nitrosated or nitrosylated PDE inhibitor of claim 1.